

THIS IS NOT A TEST REQUEST FORM. Please complete and submit with the test request form or electronic packing list.

PEDIATRIC/ADULT CYTOGENETIC TESTING PATIENT HISTORY FORM

Patient Name: _____ **Date of Birth:** _____
Sex Assigned at Birth: Female Male Intersex **Gender Identity (optional):** Female Male _____
Ordering Provider: _____ **Provider's Phone:** _____
Practice Specialty: _____ **Provider's Fax:** _____
Genetic Counselor: _____ **Counselor's Phone:** _____

Sample Type: Peripheral blood Cord blood Buccal Skin Biopsy

Desired Testing

- | | |
|---|---|
| <input type="checkbox"/> 2002289 Chromosome analysis, peripheral blood
<input type="checkbox"/> 2002287 Chromosome analysis, rule out mosaicism
<input type="checkbox"/> 2003414 Cytogenomic SNP microarray, blood
<input type="checkbox"/> 2006267 Cytogenomic SNP microarray, buccal swab
<input type="checkbox"/> 2005763 Chromosomes with reflex to microarray | <input type="checkbox"/> 2002286 Chromosome analysis, skin biopsy
<input type="checkbox"/> 0040208 Newborn FISH panel (13, 18, 21, X, Y)
<input type="checkbox"/> 2002299 FISH for specific condition (specify probe or condition): _____
_____ |
|---|---|

Indication for Testing (check all that apply—*required*)

- Suspected diagnosis of: Down syndrome Trisomy 18 Trisomy 13 Turner syndrome Other: _____
- High risk cfDNA screen (NIPT): T21 T18 T13 45,X XXX XXY XYY Other: _____
- Atypical cfDNA screen (NIPT) for chromosome: 13 18 21 X Y Other: _____ CNV Mosaicism
- Maternal** finding by cfDNA for chromosome: 13 18 21 X Y Other: _____ CNV Mosaicism
- Congenital anomalies (specify): _____
- Dysmorphic features (specify): _____
- | | |
|---|--|
| <input type="checkbox"/> Cardiac defect (specify): _____

<input type="checkbox"/> Genital anomalies: _____
<input type="checkbox"/> Ambiguous genitalia
<input type="checkbox"/> Intellectual and/or developmental disability
<input type="checkbox"/> Autism/Autism spectrum disorder
<input type="checkbox"/> Learning disabilities | <input type="checkbox"/> Infertility
<input type="checkbox"/> Recurrent miscarriage
<input type="checkbox"/> Partner with recurrent miscarriage (partner's name): _____
<input type="checkbox"/> Other (specify): _____

<input type="checkbox"/> Family history (complete information in box below) |
|---|--|

There is a family history of a chromosome or microarray abnormality in patient's (relationship to patient): _____

Confirm an abnormality *previously identified in this patient*.

IF EITHER OF THE ABOVE IS TRUE, to ensure correct testing, please provide:

1. The name/DOB of the previously tested family member (if not patient): _____
2. The abnormality found in the patient or family member: _____
3. **A copy of the family member's/patient's previous test results.**

For questions, contact an ARUP genetic counselor at 800-242-2787 ext. 5222.

Master Label